

ADHESIVE LABEL WITH GRID FOR MICROSCOPE SLIDE

5 The present invention relates to a microscope slide in combination with a self-adhesive label for use in manual microarraying.

10 Microarraying involves the transfer of e.g. biological DNA material from a source e.g. a microtitre plate to a target e.g. a glass microscope or microarray slide. Microtitre plates holding 96, 384 or 1534 different DNA samples are known, and conventionally it is desired to accurately transfer the samples from the source onto the target in the form of small distinct and
15 separate micro spots. Once the material has been deposited in the form of micro spots the individual samples can then be analysed.

20 The accurate transfer of material from a source to a target can be achieved by either robotic or manual spotting. For robotic transfer the configuration of the material on the source can be easily monitored and controlled by setting the robotic system to spot in pre-defined areas and in a pre-defined configuration.

25 However, for manual transfer of material, the placement of the material from the source onto the target tends to be more difficult to monitor and control. The material being transferred is often clear and thus once it is dry it can be very difficult to know
30 where a spot has already been placed and therefore exactly where the next spot should be placed on the target.

 It is therefore desired to overcome the problems associated with conventional manual spotting techniques.

35 According to a first aspect of the present invention there is provided the combination of a microscope slide and an adhesive label as claimed in claim 1.

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According to a second aspect of the present invention there is provided an adhesive label as claimed in claim 5.

Sub B1 5 } According to a third aspect of the present invention there is provided a method as claimed in claim 6.

According to a fourth aspect of the present invention there is provided a method as claimed in claim 7.

10 According to a preferred embodiment a pre-gridded microscope slide is provided which has a pre-printed alpha-numeric grid attached to the underside of the slide. The alpha-numeric grid appears through the glass slide with the numbers and letters in the correct
15 configuration i.e. write read up.

Various embodiments of the present invention will now be described, by way of example only, and with reference to the accompanying drawings in which:

20 Fig. 1 shows a microscope slide with a label attached thereto; and

Figs. 2(a) and 2(b) show two different labels.

With reference to Fig. 1, transfer of material onto a target microscope slide 1 is carried out from a source plate to pre-defined positions on the top surface of the
25 glass slide 1, using a pre-printed grid 2, preferably printed on a label 3, as a guide for placement of material to be spotted. Knowing which box or cell of the grid 2 has previously been spotted enables a user to safely spot the next sample in the next available empty
30 box or cell of the grid or array 2.

The format of the pre-printed grid 2 may vary, but a 8 x 12 array or grid corresponding with a 96 well microtitre plate format is preferred. According to such an embodiment, 96 different DNA samples can be
35 transferred from the source microtitre plate to the target microscope slide 1 with a one to one correspondence between the two of them.

Two different embodiments of label design are shown

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in Figs. 2(a) and 2(b). The labels are shown enlarged. In the embodiment shown in Fig. 2(a) each array 2 is approximately 6 mm x 4 mm i.e. each cell is approximately 0.5 mm x 0.5 mm. In the embodiment shown
5 in Fig. 2(b) each array 2 is approximately 12 mm x 8 mm i.e. each cell is approximately 1 mm x 1 mm. The size and number of grids 2 on a single microscope slide 1 may vary depending on the amount of material that needs to be transferred. For the transfer of relatively large
10 sample amounts, the grids 2 can be made correspondingly larger so that the individual cells of the grid 2 can hold sufficient material.

After spotting, the pre-gridded slides 1 can be used in further analysis stages which may require the
15 slides 1 to withstand extreme temperatures. The pre-gridded slides 1 can preferably withstand repeated freezing, including temperatures down to -20°C and repeated heating, up to 96°C over prolonged periods of time. The slides 1 are also preferably resistant to
20 corrosive chemicals and reagents.

The grids 2 have also been shown not to interfere with scanning of the glass slides 1 to detect
fluorescent dyes, which is usually one of the final stages of spotted material analysis.

25 Preferably, the printed grids can be removed from the glass microscope/arraying slides 1, after transfer of material has occurred.